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(54) **LOCKABLE SAFETY SHIELD ASSEMBLY FOR A PREFILLABLE SYRINGE**

VERRIEGELBARE SCHUTZSCHILDANORDNUNG FÜR EINE VORFÜLLBARE SPRITZE

**DISPOSITIF DE PROTECTION VERROUILLABLE POUR SERINGUE POUVANT ETRE
PREALABLEMENT REMPLIE**

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EP 0 941 134 B1

Description

I. Field of the Invention.

[0001] The invention relates to a lockable safety shield assembly for a prefillable syringe, and more particularly, to a lockable safety shield assembly for a prefillable syringe which is easily attached to a prefillable syringe and which can be subsequently activated by an end-user to form a locking mechanism for the shield.

II. Background.

[0002] As is known in the art, syringes are medical delivery devices utilizable to administer a medicament to a patient. Syringes are normally provided in prefillable form, wherein a set dosage of medicament can be filled into the syringe by a pharmaceutical manufacturer for distribution to the end user, or they are empty and intended to be filled by the end user from a vial or other source of medicament at the time administration of the medicament is desired.

[0003] Syringes typically include a barrel portion adapted to retain the medicament. The distal end of the barrel is normally configured to mate with a conventional piercing element, such as a pointed needle cannula made of steel or like material or a blunt ended cannula formed of plastic, to deliver the medicament contained in the barrel. A plunger rod is inserted through the open distal end of the syringe barrel and, through its engagement with an elastomeric or rubber-like stopper element fitted in a fluid-tight manner within the interior of the barrel, a user can apply manual force to the plunger to deliver the medicament through the piercing element.

[0004] More typically the case with pointed needle cannulae, accidental needle sticks that occur after the needle cannula has been used generally pose a greater health risk. To avoid such accidents, many prior art hypodermic syringes include a rigid cylindrical safety shield telescoped over the syringe barrel. The prior art safety shield can be slid from a proximal position where the needle cannula is exposed for use, to a distal position where the safety shield protectively surrounds the needle cannula.

[0005] Most prior art hypodermic syringes with cylindrical safety shields telescoped over the syringe barrel include structure for locking the safety shield in its distal position. Examples of such structures are found, for instance, in U.S. Patent Nos. 5,342,309 and 5,385,555 to Hausser and U.S. Patent No. 5,304,149 to Morigi.

[0006] While generally suitable for the tasks employed, further improvements are still being sought. For instance, where the syringe is prefillable by a pharmaceutical manufacturer, it would be beneficial to incorporate improvements into the locking structure employed with the safety shields to better facilitate a pharmaceutical manufacturer's processing of the various components associated with the syringe barrel and safety

shield. These improvements should also facilitate the end user's normal use of the prefillable syringe and easy activation of the safety shield once safe disposal of the syringe is desired. Moreover, most of the prior art approaches focus their efforts on employing a plastic collar element on some distal portion of the syringe barrel, with the shield locking onto the collar. While a valid approach, because many of the prefilled syringes currently in use are made of glass, it can be difficult to assemble the plastic collar element onto the syringe barrel.

[0007] According to US 5,697,908, a lockable safety shield for a prefillable syringe is provided. The design of the lockable safety shield enhances pharmaceutical manufacturers' ease of assembling the various components as part of its filling or processing of the prefillable syringes in normal practice. A collar is attachable to the distal hub of the prefillable syringe. A safety shield is axially slidable over the collar between a retracted position, wherein the distal end of the piercing element associated with the prefillable syringe is exposed, and an extended position, wherein the safety shield is locked to the collar to protectively cover the distal end of the piercing element. The safety shield includes locking structure configured so that the shield, as supplied to the pharmaceutical manufacturer, can be easily fitted to the syringe. The locking structure includes at least one deflectable arm provided on the body of the shield. The deflectable arm includes a proximal end deflectable towards the interior of the shield. A stop member is provided on the interior of a shield in spaced relation to the proximal end of the deflectable arm. A ring is axially slidable over the shield to deflect the arm towards the interior of the shield to activate the locking structure. The safety shield is slid distally by an end user such that the collar is lockingly retained between the stop member and the proximal end of the deflectable arm. The collar, the stop member, and the deflectable arm may be configured to provide tactile as well as audible indication of locking to the end user.

[0008] EP-A-740,942 relates to a method of manufacturing a pre-filled injection device, comprising the steps of forming, from a deformable plastics material, a tubular syringe body provided with an external blocking member capable of deforming radially between retracted and extended positions, of forming tubular sheath provided with an internal blocking member interacting with that of the syringe body, a mounting the sheath around the syringe body by the relative sliding thereof, rendered possible by the radial retracting of the blocking member of the syringe body, and of introducing, into the syringe body, a pre-filled rigid container capable of moving the blocking member of the said syringe body into its extended position after injection, thus irreversibly securing the sheath in an advanced position of protection after use once the said blocking member has been moved.

III SUMMARY OF THE INVENTION

[0009] According to the present invention, there is provided a medical device as defined in claim 1. **IV. Details of the Invention.**

[0010] A lockable safety shield assembly for prefillable syringes is provided. The lockable safety shield is designed to facilitate a pharmaceutical manufacturer's assembly of the various components associated with the lockable safety shield onto the syringe in conjunction with how the syringes are normally processed by the pharmaceutical manufacturer. The locking structure associated with the safety shield of the present invention may be designed such that it will not interfere with easy assembly by the pharmaceutical manufacturer, but which can be easily activated by an end user to facilitate safe covering of the distal end of a piercing element once the prefillable syringe has been used.

[0011] The assembly includes a tube mounted about the outside surface of the syringe barrel. The tube can be locked, for instance, to the flange provided about the proximal end of the syringe barrel. The tube features a collar positioned adjacent the distal end of the syringe barrel. The collar can be configured as a flat, disk-like structure formed in an annular manner about the outside of the tube.

[0012] A safety shield is coaxially mounted over the tube. The safety shield is axially movable between a retracted position, wherein the distal tip of a piercing element affixable to the barrel is exposed, and an extended position, wherein the distal tip of the piercing element will be covered. The safety shield includes a proximal end, a distal end, and a circumferential sidewall therebetween. The circumferential sidewall has an outside surface and an inside surface. Particularly, the shield is provided with user-activatable locking structure to secure the shield in its extended position relative of the syringe.

[0013] The locking structure includes at least one deflectable arm incorporated as part of the structure of the sidewall. The deflectable arm includes a distal end mounted with the circumferential sidewall, and a proximal end which is deflectable towards the interior of the shield. At least one stop member is mounted to the interior surface of the shield and spaced a distance from the proximal end of the deflectable arm to define a gap between the stop member and the proximal end of the deflectable arm. The deflectable arm has a free position, wherein the outside surface of the arm is substantially co-planar with the outside surface of the circumferential sidewall, and a locking position, wherein the deflectable arm is urged towards the interior of the shield so that the proximal end of the deflectable arm is positioned towards the interior of the shield.

[0014] Means are provided on the shield to activate the locking structure. A circumferential ring is axially slidable about the outside surface of the shield between a first position, wherein the ring is spaced away from the

deflectable arm, and a second position, wherein the ring is positioned against the at least one deflectable arm to secure the arm in the locking position. The ring may be secured to the distal end of the shield by a user-severable connection. The user-severable connection may entail, for instance, one or more frangible sections of material provided between the ring and the distal end of the shield. A pair of spaced abutments may be provided on the outside surface of the deflectable arm to retain the ring in its second position.

[0015] In practice, the safety shield assembly is typically preassembled by the syringe manufacturer and supplied to the pharmaceutical manufacturer in a sterile state, ready for attachment to the prefillable syringe once it has been filled with a desired drug.

[0016] After the syringe has been filled with the drug and otherwise subjected to various processing operations, such as various particulate inspection operations, labeling operations, plunger rod assembly operations, or the like, the safety shield assembly can be easily fitted to the syringe. The tube is fitted about the syringe barrel and clipped onto the flange of the syringe barrel. In the preassembled state, the shield located relative of the tube such that the shield will be placed in a retracted position vis-a-vis the piercing element. Various structures associated with the tube may be provided with sloped surfaces to ensure smooth placement of the safety shield assembly over the syringe flange.

[0017] Subsequent to placing the shield assembly onto the syringe barrel, the locking structure must be activated. The locking structure is activated by detaching the ring from its user-severable connection, and sliding the ring proximally so that it is engaged between the abutments provided on the deflectable arm. If desired, the device may be shipped by a pharmaceutical manufacturer to an end user with the ring connected to the shield by its user-severable connection, so that the end-user himself activates the locking structure. Alternately and preferably, the pharmaceutical manufacturer activates the locking structure as part of the assembly process, such that the filled syringe with safety shield assembly is shipped to an end-user with the locking structure already activated.

[0018] After the syringe has been used to administer an injection, the shield is slid distally over the tube. The collar will be captured between the proximal end of the deflectable arm and the stop member in order to lock the shield in its extended position. In a preferred embodiment, the deflectable arm, the collar, and the stop member may be configured to provide audible as well as tactile indication to the end user that the shield has been securely locked with the collar.

[0019] In a preferred embodiment, at least two deflectable arms are provided. The two deflectable arms are located on diametrically opposite sides of the shield. Correspondingly, two stop members are provided to interact with the two deflectable arms. When viewed along the central axis of the shield, the stop members can be

circumferentially aligned with a respective proximal end of the deflectable arms, or they can be circumferentially offset from a respective proximal end.

V. Brief Description of the Drawings

[0020] The invention will now be described in greater detail by way of reference to the appended drawings, wherein Figures 13 to 16 illustrate the present invention and the remaining Figures 1 to 12 are retained as an aid to understanding:

Figure 1 depicts, in perspective view, a lockable safety shield assembly mounted to a syringe;

Figure 2 is an exploded perspective view of the lockable safety shield assembly of Figure 1;

Figure 3 is a cut-away view of the lockable safety shield being mounted over the tube and the syringe barrel;

Figure 4 is a cross-sectional view of the lockable safety shield assembly subsequent to mounting to the syringe;

Figure 5 is a cross-sectional view of the lockable safety shield assembly after an end user has activated the locking mechanism;

Figure 6 is a cross-sectional view of the lockable safety shield assembly after it has been urged in a distal direction by an end-user to protectively cover the needle cannula;

Figure 7 is a partial cross-sectional view of the lockable safety shield assembly illustrating activation of the locking mechanism;

Figure 8 is a partial cross-sectional view of the lockable safety shield assembly interacting with the collar located on the tube;

Figure 9 is an end elevational view of the lockable safety shield assembly;

Figure 10 is a partial cross-sectional view of a manner of configuring the collar;

Figure 11 is a cross-sectional view of an alternative embodiment of the lockable safety shield assembly after it has been urged in a distal direction by an end-user to protectively cover the needle cannula;

Figure 12 is an enlarged, partial cross-sectional view of the alternative embodiment of the lockable safety shield assembly illustrated in Figure 11 interacting with the collar located on the tube;

Figure 13 depicts, in perspective, a partial cross-sectional view of an embodiment of the lockable safety shield assembly of the present invention mounted to a syringe;

Figure 14 is a fragmentary side view of the flange portion of the an embodiment of the lockable safety shield assembly illustrated in Figure 13;

Figure 15 is a fragmentary planar view of the flange portion of an alternative embodiment of the lockable safety shield assembly illustrated in Figure 13; and Figure 16 depicts a fragmentary schematic repre-

sentation of the insertion of the syringe into the flange portion of the lockable safety shield assembly illustrated in Figure 13.

VI. Detailed Description of the Invention

[0021] A convention utilized throughout this description is that the term "distal" refers to the direction furthest from a practitioner, while the term "proximal" refers to the direction closest to a practitioner.

[0022] Turning now to the drawings, wherein like numerals denote like components, Figures 1-4 depict one embodiment of a lockable safety shield assembly 30. The lockable safety shield assembly 30 can be provided in conjunction with a prefillable syringe 10. As illustrated in Figs. 1 and 2, prefillable syringe 10 includes a barrel 16 characterized by a proximal end 14, a distal end 12, and a hub 24 provided adjacent the distal end. A flange 18 may be provided about the proximal end, with a plunger rod 20 inserted through proximal end 14. Plunger rod 20 is attached to a stopper 22 used for urging fluid through barrel 16. Fluid is emitted or aspirated via a fluid path, structured at distal end 12 of the syringe, that communicates with interior portions of barrel 16. For instance, if the pre-fillable syringe is configured as a luer lock syringe, the fluid path is configured as a luer tip (not shown) provided at distal end 12. Here, pre-fillable syringe 10 is illustrated with a fluid path configured as a piercing element 26, such as a pointed needle cannula. Piercing element 26 is characterized by a distal end 28.

[0023] The lockable safety shield assembly 30 includes a tube 200 which is mounted about barrel 16 of the syringe. The tube 200 features a proximal end 204, a distal end 202, and a circumferential sidewall 206 therebetween. A flange structure 208 is provided adjacent proximal end 204 of the tube. As best seen in Figure 3, flange structure 208 includes a lip 210 sized to lockingly engage flange 18 of the syringe barrel when tube 200 is placed over barrel 16. Adjacent distal end 202 of the tube there is provided a collar 32. Collar 32 can be formed as a disk-like structure characterized by an edge 34 and a side portion 36. Collar 32 has a thickness "e" and is characterized by a maximum diameter "g". Collar 32 may be provided as part of the structure of tube 200. Alternately, as the skilled artisan will appreciate, collar 32 may be separately supplied and retained to the tube by friction fit, adhesives, mechanical means, or like structure. If desired, edge 34 can display one or more raised nubs 39 having grooves 40. Grooves 40 can be aligned with complementary rails (not shown) provided on an interior portion of safety shield 42 for aligning the safety shield during the assembly process.

[0024] For purposes of clarity, Figures 1-6 depict collar 32 as having a rectangular cross-section, with side portion 36 displaying relatively flat walls that are directly linked to edge 34. If so desired, collar 32 can be configured in this manner to construct a lockable safety shield assembly. However, Figure 10 depicts a preferred way

for configuring the collar (here designated by the numeral 32a) so to provide better audible as well as tactile indication that shield 42 has been locked to the collar, as well as to enhance locking action between the shield and the collar, as will be further explained.

[0025] Referring then to Figure 10, as before, collar 32a includes an edge 34 and a side portion 36. However, here, side portion 36 is not completely flat. Rather, when viewed in cross-section, edge 34 and side portion 36 are linked by a locking tooth 34a. Locking tooth 34a features a distally-facing canted edge 36a and a proximally-facing canted edge 36b. Each of canted edges 36a, 36b of the locking tooth terminate at a shelf 37 that intersects with side portion 36. As before, if desired, grooves (not shown) can be provided on edge 34, for the purposes previously explained. As before, collar 32a includes a thickness "e" and a maximum diameter "g". Locking tooth 34a includes a maximum width "q" which need not be equal to thickness "e" displayed by collar 32a.

[0026] Safety shield 42 is characterized by a proximal end 44, a distal end 45, and a circumferential sidewall 46 therebetween. Circumferential sidewall 46 includes an outside surface 48 and an inside surface 49. Outside surface 48 is characterized by a diameter "N". Shield 42 is slidable both over tube 200 and collar 32, and thus over syringe barrel 16, between a retracted position (Figs. 1, 4 and 5), wherein distal tip 28 of piercing element 26 is exposed, and an extended position (Fig. 6), wherein the distal tip of the piercing element is protectively covered.

[0027] Lockable safety shield assembly 30 includes user-activatable locking structure. A feature of the lockable safety shield assembly is that the locking structure can be implemented such that it is not "activated" during assembly of the components, making it easier to assemble the components to syringe 10. Subsequent to assembly of the components, the locking structure can be "activated", putting the safety shield assembly in a ready-to-use state.

[0028] Making reference to Figs. 1-3 and 7-9, shield 42 features at least one deflectable arm 50. Arm 50 is preferably provided as part of the structure of the shield. Arm 50, which may be molded as part of the shield or thereafter created from the shield via various cutting processes, includes a distal end 54 and a proximal end 52. Distal end 54 is resiliently affixed to sidewall 46 such as by a living hinge arrangement, while proximal end 52 is not attached to the sidewall and is configured for free movement towards an interior portion of shield 42. In a preferred embodiment, at least two deflectable arms 50 are provided on shield 42, with these two arms 50 located on diametrically-opposite sides of shield 42. Arms 50 are thus configured for a free position (Figures 1 and 4), wherein the surface of the arms are substantially coplanar with circumferential sidewall 46 of the shield, and a locking position (Figures 5, 7, and 8), wherein proximal ends 52 of arms 50 are urged towards the interior of the shield.

[0029] As best reflected by Figures 7 and 8, in a preferred configuration, proximal ends 52 of the deflectable arms are not straight. Rather, to best make use of the features of collar 32a (Figure 10), proximal ends 52 are preferably canted in a manner such that when the deflectable arms are positioned towards the interior of the shield, the proximal ends will be oriented to lockingly cooperate with distally-facing canted edge 36a of collar 32a, as will be explained hereinbelow.

[0030] At least one stop member 70 is provided on inside surface 49 of the shield in an area adjacent proximal end 44. In a preferred embodiment, two stop members 70 are provided, each corresponding to a respective one of two deflectable arms 50 (see Fig. 9). Stop member 70 includes a distally-facing stop surface 72. Stop surface 72 is preferably spaced from proximal end 52 of the deflectable arm a distance "b" as measured along central axis "z" of the shield (see Fig. 3).

[0031] As best illustrated in Figures 7 and 8, in a preferred configuration, stop surface 72 is preferably not straight. Rather, to best make use of the features of collar 32a (Figure 10), stop surface 72 is preferably canted in a manner such that it will lockingly cooperate with proximally-facing canted edge 36b of collar 32a, as will be explained hereinbelow.

[0032] Stop member 70 also includes a proximally-facing sloped surface 74. Proximally-facing sloped surface 74 facilitates sliding of shield 42 over collar 32 when the shield is assembled to tube 200. Stop member 70 can be located such that stop surfaces 72 are circumferentially aligned with respective proximal ends 52 of deflectable arms 50, when viewed along central axis "Z" of the shield. Alternately, if desired, stop members 70 can be placed on interior surface 49 of the shield such that they are circumferentially offset from respective proximal ends 52 of the deflectable arms, when viewed along central axis "Z" of the shield.

[0033] Shield 42 includes structure permitting activation of the deflectable arms 50 into their locked position (Figs. 5, 6, 7, 8). A respective pair of distal and proximal abutments 58, 56 are provided on deflectable arms 50. Distal abutment 58, which includes a distally-facing sloped surface 60, is spaced from proximal abutment 56 by a distance "a" that defines a gap 59. As seen in Figs. 3 and 4, in a preferred embodiment where two diametrically opposite deflectable arms are included, the diametrically opposite pair of gaps 59 define a diameter "M" slightly greater than diameter "N" defined by outside surface 48 of the shield when the arms 50 are in their free position. The effect is that gap 59 is configured so that it is elevated from the level of surrounding outside surface 48 of cylindrical sidewall 46 prior to activation of deflectable arm 50.

[0034] A circumferential ring 80 having an inside wall 82 is disposed for coaxial movement with circumferential sidewall 46 of the shield. Inside wall 82 has a diameter "P" at least equal to, if not greater than, diameter "N" defined by the outside surface of shield 42. Ring 80

has a width "d" at least equal to, if not slightly less than, the width of gap 59 as defined by distance "a". To simplify later assembly, shield 42 is typically configured with ring 80. Ring 80 can be retained to shield 42, for instance, by a user-severable connection. That is to say, shield 42 and ring 80 are preferably supplied to the pharmaceutical manufacturer as a single unit. For example, ring 80 can be supplied by the component manufacturer such that it is secured to the shield by a user-severable connection such as a threaded connection. Preferably, and as is disclosed herein, ring 82 is secured to distal end 45 of the shield by one or more user-severable connections 84 configured as thinned, frangible sections of material between ring 80 and proximal end 45 of the shield, thereby permitting easy removal of the ring from the shield for further assembly operations. This type of structure also permits the component manufacturer to easily form ring 80 and shield 42 as a single molding.

[0035] Shield 42 further includes structure for retaining shield 42 in the retracted position without being inadvertently "over-retracted" with respect to syringe 10 and, hence, inadvertently removed from syringe 10 in the proximal direction. As seen in Fig. 3, one or more distal stop members 76 can be provided on interior surface 49 of the shield adjacent distal end 45. Distal stop members 76 are configured to engage a distally-facing side portion 36 of the collar to prevent shield 42 from being over-retracted in a proximal direction. If desired, one or more tertiary stop members 78 can be provided on shield 42, each spaced from a respective distal stop member 76 by a distance "c" approximately equal to thickness "e" of collar 32. Hence, collar 32 can be releasably retained between distal stop members 76 and tertiary stop members 78 when the shield is in its retracted position. If desired, tertiary stop member 78 can be provided with a proximally-facing sloped surface 79 which, like proximally-facing sloped surface 74 associated with stop members 70, assists in the assembly of shield 42 about collar 32.

[0036] Operation of a device will now be explained.

[0037] As seen in Figure 3, means can be fitted to the syringe adjacent the distal end to provide a barrier to the fluid path leading to the medicament which will be contained within barrel 16. The barrier is normally fitted by the syringe manufacturer and here, can be fitted to the syringe barrel in a process subsequent to placement of collar 32. If the fluid path is structured as a luer tip, for instance, a luer tip cap can be fitted. Here, where a piercing element 26 is configured on the syringe, a well-known needle shield 29 can be fitted about piercing element 26 so that a proximal-end of needle shield 29 rests against collar 32. Collar 32 should be configured such that maximum diameter "g" is at least slightly greater than a maximum diameter "f" displayed by needle shield 29, such that needle shield 29 will not interfere between operation of collar 32 and shield 42.

[0038] Preferably, safety shield assembly 30 is preassembled by the component manufacturer and present-

ed to the pharmaceutical company in a sterile state, ready to be fit to the syringe once filled with the drug. Syringe 10 is supplied to the pharmaceutical manufacturer in a sterile state, ready to be filled with a desired drug and otherwise processed in a well-known manner. For instance, various filling operations, particulate inspection operations, labeling operations, plunger rod assembly operations may be conducted on the syringe.

[0039] As a first step in assembling the safety shield assembly, shield 42 is fitted to tube 200. Proximally-facing sloped surfaces 74 and, if provided, 79, assist shield 42 to smoothly surpass edge 34 of the collar. Shield 42 is slid over tube 200 in a proximal direction, such that collar 32 rests against distal stop members 76. If tertiary stop members 78 are provided, collar 32 will be releasably retained between distal stop members 76 and tertiary members 78. After the various filling and processing operations have been conducted on the syringe, tube 200 is slid over syringe barrel 16 such that flange 18 of the syringe is captured by lip structure 210 of the tube. As the shield is held to the tube in a retracted position, with needle shield 29 affixed to piercing element 26 will project beyond the distal end 202 of the tube. Syringe 10, together with lockable safety shield assembly 30, can be shipped by the pharmaceutical manufacturer to an end-user in a ready-to-use state.

[0040] Fig. 4 is representative of safety shield 42 mounted about syringe 10 prior to activation of locking structure, while Fig. 5 is representative of the same construction after activation of the locking structure. Preferably, a pharmaceutical manufacturer will ship the prefillable syringe to the end-user in the state shown in Fig. 5, such that the locking structure is pre-activated. One way to do this is that the locking structure may be activated by the component manufacturer, as a final step in the preassembly of the safety shield assembly 30. For instance, the step of sliding ring 80 towards gap 59 can be effected as a final step in an assembly motion during which shield 42 is fitted to tube 200. In this vein, it will also be realized that because the component manufacturer preferably supplies the ring and shield to pharmaceutical manufacturer as a preassembled, single unit, the later assembly operations by the pharmaceutical manufacturer are simplified, in that the pharmaceutical manufacturer need only be concerned with sorting, orienting, distributing and then assembling a single unit about the syringe. Alternately, of course, it will be realized that the activation step can be done by the pharmaceutical manufacturer itself, as part of the operation in fitting the safety shield assembly to the syringe.

[0041] Nonetheless, Fig. 5 illustrates activation of the locking structure, whether by the component manufacturer, by the pharmaceutical manufacturer or by an end-user. The user-severable connections 84 connecting ring 80 to distal end 45 of the shield have been severed, and ring 80 has been slid proximally with respect to shield 42. Aided by distally-facing sloped surface 60 associated with distal abutment 58, inside surface 82 of

the ring glides along outside surface 48 of the shield, over distally-facing sloped surface 60, and is secured in gap 59 defined between distal abutment 58 and proximal abutment 56. Because inside diameter "P" of ring 80 is smaller than diameter "M" between diametrically-opposed gaps 59, ring 80 exerts an inwardly-directed force upon deflectable arms 50. Deflectable arms 50 are thus deflected about their distal ends 54, such that proximal ends 52 are pushed towards the interior of shield 42. As a deflectable arm 50 is urged inwardly (see particularly Figs. 7 and 8), proximal end 52 of a given deflectable arm and stop surface 72 of a respective stop element 70 define a gap 100 between them. Gap 100 is designed to "capture" collar 32 when the shield is slid distally to cover piercing element 26. Where the collar has a flat side portion 36, then gap 100 should have a width "b" at least equal, if not slightly greater than, thickness "e" of the collar to securely retain collar 32 in gap 100. If, as depicted in Figure 10, the collar is configured with locking tooth 34a, then width "b" should be chosen so as to accommodate the maximum width "q" associated with locking tooth 34a, so that locking tooth 34a is securely retained in gap 100.

[0042] Fig. 6 is representative of shield 42 in its extended position to protectively cover distal tip 28 of the piercing element. After needle shield 29 has been removed and an injection administered in the usual manner, shield 42 is grasped by an end-user and slid distally with respect to syringe 10. Collar 32, which is fixed to tube 200, will thus be urged towards proximal end 44 of the shield. Owing to the resiliency of deflectable arms 50, edge 34 of the collar will push arms 50 in an outward direction from the interior of the shield and thus will pass along the length of arms 50. Edge 34 will eventually deflect past proximal end 52, causing the arm to re-deflect in an inward direction toward the interior of the shield. An audible indication, such as a clicking sound, will be heard to indicate activation of the safety mechanism. Thus, collar 32 will come to rest in gap 100, with the collar sandwiched between proximal end 52 and stop surface 72. Collar 32 is thus lockingly held between stop surface 72 and proximal end 52 of the arm, arresting further movement of the shield in either proximal or distal directions respective of syringe 10. Accordingly, shield 42 is retained in its extended position with respect to piercing element 26, protectively covering distal tip 28 against inadvertent touch contact.

[0043] As previously explained, in a preferred configuration (Figures 7, 8 and 10), deflectable arms 50, stop members 70 and collar 32a are configured to provide good audible as well as tactile indication that shield 42 has been locked to collar 32a. As edge 34 of collar 32a clears proximal end 52 of the deflectable arm, locking tooth 34a enters gap 100. As soon as the locking tooth enters gap 100, the resilient characteristics of the deflectable arm cause the arm to re-deflect in an inward direction toward the interior of the shield. At this moment, an interior portion 49a of deflectable arm 50 that

is located adjacent proximal end 52 is thrust against shelf 37 of collar 32a. As soon as interior portion 49a is thrust against shelf 37, tactile as well as audible sensations are transmitted to the user, signaling that shield 42 has been locked to collar 32a. Owing to the canted orientations of both proximal end 52 of the deflectable arms and stop surface 72 of the stop members 70, each of the proximal ends 52 and stop surfaces 72 are lockingly retained against their respective distally facing canted edge 36a and proximally-facing canted edge 36b, such that tooth 34a is securely locked within gap 100. Thus, by a simple distal movement, shield 42 is self-locking with collar 34a.

[0044] Referring then to Figure 11, and Figure 12 for greater detail, an alternative preferred embodiment of the collar 132 is illustrated, with the collar including an edge 134 and a side portion 136. However, here, side portion 136 is not completely flat. Rather, when viewed in cross-section, edge 134 and side portion 136 are not linked by a locking tooth. Instead, the side portion 136 includes a substantially flat abutting surface 136a and a distally-facing sloped surface 138. In this way, as proximal end 52 of the deflectable arm 50 is thrust towards the tube it abuts surface 136a of collar 132. Owing to the canted orientations of both proximal end 52 of the deflectable arms and sloped surface 138, if safety shield 42 is thrust against the tube 200, the proximal end 52 each will travel along sloped surface 138 towards the syringe 10 thereby preventing exposure of the needle.

[0045] In a preferred embodiment of this invention, as shown in Figures 13 through 16, the flange 208 of the tube 200, includes an upper proximal collar 210 spaced apart from said flange and supported on the flange by a plurality of posts 212. Preferably, the collar 210 has a polygonal shape, with an inwardly-sloping edge 214 along its inner surface and terminating a shoulder 218. In this way, as the syringe 10 is inserted into the tube 200, the flange 18 of the syringe comes in contact with the edge 214 causing the collar 210 to flex as the flange travels along the ridge. Once past the shoulder 218, the collar 210 is resilient enough to return to its resting position with the leading edge 218a of the shoulder extending beyond point at which the curved surface of the flange 18 transitions to the planar surface TP. In this way, the leading edge 218a will not readily come in contact with the curved portion of the flange so that the syringe cannot be easily removed from safety shield assembly 30.

[0046] The various components can be formed from suitable materials, such as medical-grade plastics or the like. They may be injection molded in manners well-known to the skilled artisan, resulting in reliable and cost-effective production of the locking shield assembly. The locking shield assembly of the present invention can be adapted with prefilled syringes of various dimensions and capacities. Various enhancements can be provided to permit easy assembly. For instance, sections of the sidewall adjacent the proximal end of the

shield can be thinned, or slits can be incorporated in an area of the shield between the proximal end of the shield and the proximal edge of the deflectable arms, all to enhance the easier sliding of the shield proximally over the collar. Of course, the dimensions for the various components can be selected such that when the shield is in its retracted position, the distal tip of the piercing element is exposable for an injection, and when the shield in its extended position, the distal tip of the piercing element will be protectively covered to enable the end user to safely discard the syringe.

[0047] It will be appreciated and understood by those skilled in the art that further and additional revisions to the invention may be devised. The invention is not limited to the specific embodiments shown.

Claims

1. A medical device comprising:

a barrel (16) having proximal (14) and distal (12) ends;
a tube (200) secured about the barrel (16);

wherein the tube (200) includes a tube flange (208);

a shield (42) mounted about the tube (200) and axially moveable between a retracted position, wherein the distal end (12) of the barrel (16) is exposed, and an extended position, wherein the distal end (12) of the barrel (16) is covered, the shield (42) having a proximal end (44), a distal end (45), and a circumferential sidewall (46) there between, the circumferential sidewall (46) having an outside surface (48), an inside surface (49) and an interior; and
a user-activatable locking means to secure the shield (42) in the extended position, the user activatable locking means including:

at least one deflectable arm (50) having a distal end (54) mounted within the circumferential sidewall (46) and a proximal end (52) deflectable toward the interior of the shield (42), the arm (50) having at free position in which the arm (50) is substantially coplanar with the circumferential sidewall (46) of the shield (42), and a locking position, wherein the arm (50) is deflected toward the interior of the shield (42) so that the proximal end (52) is positioned toward the interior of the shield (42);

at least one stop member (70) mounted to the interior surface (49) of the shield (42) adjacent to the proximal end (44) of the shield (42) to define a gap between the

stop member (70) and the proximal end (52) of the deflectable arm (50); and
a ring (80) axially slidable around the outside surface of the shield (42) between a first position wherein the ring (80) is spaced away from the deflectable arm (50) and a second position wherein the ring (50) is positioned against the deflectable arm (50) to secure the arm (50) in the locking position, the tube (200) includes a collar (32) positioned adjacent the distal end (12) of the tube (200), the collar (32) having an edge (34), the tube (200) further includes an upper proximal collar (210) spaced apart from the tube flange (208) and supported on the flange (208) so that, as the barrel (16) is inserted into the tube (200), the proximal end of the barrel (16) comes in contact with the upper proximal collar (210), causing the upper proximal collar (210) to flex from its resting position, and, as the proximal end (14) of the barrel (16) passes the upper proximal collar (210), the upper proximal collar (210) is resilient enough to return to its resting position.

2. The medical device of claim 1 wherein the upper proximal collar (210) is supported on the tube (200) by a plurality of posts (212).

3. The medical device of claims 1 or 2 wherein the upper proximal collar (210) has a generally polygonal shape with an inwardly sloping edge (214) along an upper surface.

4. The medical device of claim 3 wherein the barrel (16) has a flange (18) provided about the proximal end (14) and the inwardly-sloping edge (214) of the inner surface of the upper proximal collar (210) terminates at a shoulder (218) having a leading edge (218a) extending beyond a point at which a curved surface of the barrel flange (18) transitions to a planar surface so that the leading edge will not readily come in contact with the curved portion of the barrel flange to prevent easy removal of the tube.

5. The medical device of claims 1, 2, 3 or 4 wherein the edge (134) of the collar (132) has a flat abutting surface (136a) and a distally-facing sloped surface (138) such that, when the shield (42) is thrust against the tube (200), the proximal end (52) will travel along the sloped surface (138).

Patentansprüche

1. Medizinische Vorrichtung, die folgendes umfaßt:

einen Zylinder (16) mit einem proximalen (14) und einem distalen (12) Ende,

eine um den Zylinder (16) befestigte Röhre (200),

bei der die Röhre (200) einen Röhrenflansch (208) einschließt,

einen Schutz (42), um die Röhre (200) angebracht und in Axialrichtung beweglich zwischen einer zurückgezogenen Position, in der das distale Ende (12) des Zylinders (16) freigelegt wird, und einer ausgefahrenen Position, in der das distale Ende (12) des Zylinders (16) umhüllt wird, wobei der Schutz (42) ein proximales Ende (44), ein distales Ende (45) und eine in Umfangsrichtung verlaufende Seitenwand (46) zwischen denselben hat, wobei die in Umfangsrichtung verlaufende Seitenwand (46) eine Außenfläche (48), eine Innenfläche (49) und ein Inneres hat, und

ein benutzeraktivierbares Verriegelungsmittel zum Sichern des Schutzes (42) in der ausgefahrenen Position, wobei das benutzeraktivierbare Verriegelungsmittel folgendes einschließt:

wenigstens einen ablenkbaren Arm (50) mit einem innerhalb der in Umfangsrichtung verlaufenden Seitenwand (46) angebrachten distalen Ende (54) und einem zum Inneren des Schutzes (42) hin ablenkbaren proximalen Ende (52), wobei der Arm (50) eine freie Position hat, in welcher der Arm (50) wesentlich in einer Ebene mit der in Umfangsrichtung verlaufenden Seitenwand (46) des Schutzes (42) liegt, und einer Verriegelungsposition, in welcher der Arm (50) zum Inneren des Schutzes (42) hin abgelenkt wird, so daß das proximale Ende (52) zum Inneren des Schutzes (42) hin angeordnet wird,

wenigstens ein an der Innenfläche (49) des Schutzes (42) angebrachtes Anschlagelement (70), angrenzend an das proximale Ende (44) des Schutzes (42), um einen Spalt zwischen dem Anschlagelement (70) und dem proximalen Ende (52) des ablenkbaren Arms (50) zu definieren, und einen Ring (80), der in Axialrichtung um die Außenfläche des Schutzes (42) verschoben werden kann, zwischen einer ersten Position, in welcher der Ring (80) mit Zwischenraum vom ablenkbaren Arm weg (50) angeordnet wird, und einer zweiten Position, in welcher der Ring (80) gegen den ablenkbaren Arm (50) angeordnet

wird, um den Arm (50) in der Verriegelungsposition zu sichern,

wobei die Röhre (200) einen angrenzend an das distale Ende (12) der Röhre (200) angeordneten Bund (32) einschließt, wobei der Bund (32) eine Kante (34) hat, wobei die Röhre (200) außerdem einen oberen proximalen Bund (210) einschließt, mit Zwischenraum zum Röhrenflansch (208) angeordnet und auf dem Röhrenflansch (208) getragen, so daß das proximale Ende des Zylinders (16) in Kontakt mit dem oberen proximalen Bund (210) kommt, wenn der Zylinder (16) in die Röhre (200) eingeführt wird, was bewirkt, daß sich der obere proximale Bund (210) aus seiner Ruheposition biegt, und der obere proximale Bund (210) ausreichend elastisch ist, um zu seiner Ruheposition zurückzukehren, wenn das proximale Ende (14) des Zylinders (16) den oberen proximalen Bund (210) passiert.

2. Medizinische Vorrichtung nach Anspruch 1, bei welcher der obere proximale Bund (210) durch eine Vielzahl von Stützen (212) auf der Röhre (200) getragen wird.
3. Medizinische Vorrichtung nach Anspruch 1 oder 2, bei welcher der obere proximale Bund (210) eine allgemein vieleckige Form mit einer nach innen abfallenden Kante (214) längs einer oberen Fläche hat.
4. Medizinische Vorrichtung nach Anspruch 3, bei welcher der Zylinder (16) einen um das proximale Ende (14) bereitgestellten Flansch (18) hat und die nach innen abfallende Kante (214) der Innenfläche des oberen proximalen Bundes (210) an einem Absatz (218) endet, der eine Vorderkante (218a) hat, die sich über einen Punkt hinaus erstreckt, an dem eine gekrümmte Fläche des Zylinderflanschs (18) in eine ebene Fläche übergeht, so daß die Vorderkante nicht ohne weiteres in Kontakt mit dem gewölbten Abschnitt des Zylinderflanschs kommen wird, um ein leichtes Abnehmen der Röhre zu verhindern.
5. Medizinische Vorrichtung nach einem der Ansprüche 1, 2, 3 oder 4, bei der die Kante (134) des Bundes (132) eine flache Stoßfläche (136a) und eine in distaler Richtung zeigende abgeschrägte Fläche (138) hat derart, daß sich das proximale Ende (52) längs der abgeschrägten Fläche (138) bewegen wird, wenn der Schutz (42) gegen die Röhre (200) geschoben wird.

Revendications

1. Dispositif médical comprenant:

un corps (16) pourvu d'une extrémité proximale (14) et d'une extrémité distale (12);
un tube (200) fixé autour du corps (16);

dans lequel le tube (200) est pourvu d'une bride de tube (208); 5

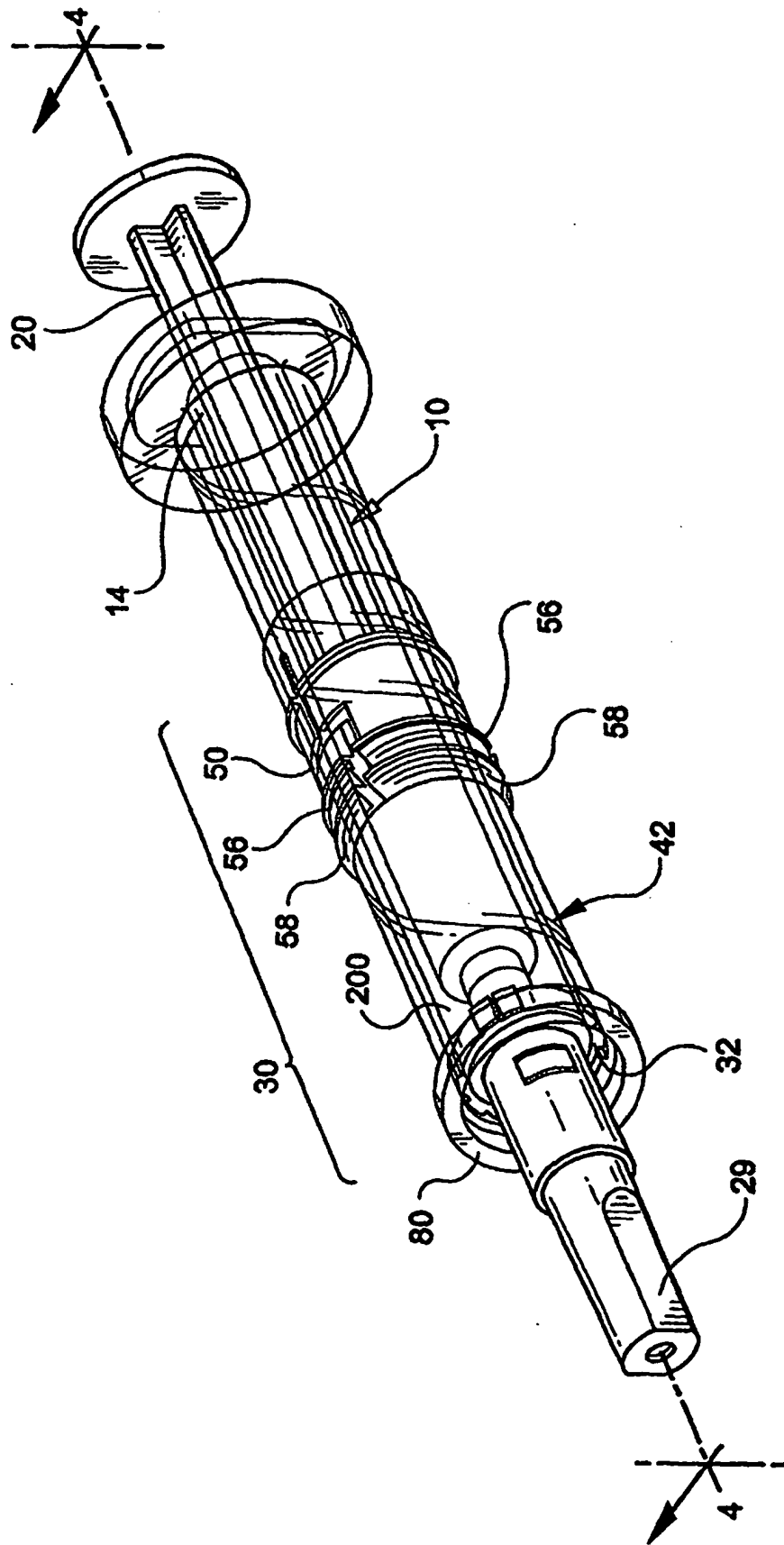
une protection (42) montée autour du tube (200) et mobile dans le sens axial entre une position rétractée, dans laquelle l'extrémité distale (12) du corps (16) est exposée, et une position étendue, dans laquelle l'extrémité distale (12) du corps (16) est couverte, la protection (42) étant pourvue d'une extrémité proximale (44), d'une extrémité distale (45) et d'une paroi latérale circonférentielle (46) entre les deux, la paroi latérale circonférentielle (46) étant pourvue d'une surface extérieure (48), d'une surface intérieure (49) et d'un intérieur; et 10
un moyen de verrouillage pouvant être activé par un utilisateur afin d'immobiliser la protection (42) dans la position étendue, le moyen de verrouillage pouvant être activé par un utilisateur incluant: 15

au moins un bras (50) pouvant être dévié pourvu d'une extrémité distale (54) montée à l'intérieur de la paroi latérale circonférentielle (46) et d'une extrémité proximale (52) pouvant être déviée vers l'intérieur de la protection (42), le bras (50) ayant une position libre dans laquelle le bras (50) est essentiellement coplanaire avec la paroi latérale circonférentielle (46) de la protection (42) et une position de verrouillage dans laquelle le bras (50) est dévié vers l'intérieur de la protection (42) de manière à ce que l'extrémité proximale (52) soit positionnée vers l'intérieur de la protection (42); 20
au moins un élément d'arrêt (70) monté sur la surface intérieure (49) de la protection (42) de manière adjacente à l'extrémité proximale (44) de la protection (42) de manière à définir un espace entre l'élément d'arrêt (70) et l'extrémité proximale (52) du bras (50) pouvant être dévié; et 25
une bague (80) pouvant coulisser dans le sens axial autour de la surface extérieure de la protection (42) entre une première position, dans laquelle la bague (80) est située à distance du bras (50) pouvant être dévié, et une seconde position, dans laquelle la bague (80) est positionnée contre le bras (50) pouvant être dévié de manière à immobiliser le bras (50) dans la position de verrouillage, 30
le tube (200) inclut un collier (32) positionné de manière adjacente à l'extrémité dis-

tales (12) du tube (200), le collier (32) étant pourvu d'un bord (34), le tube (200) inclut, en outre, un collier proximal supérieur (210) situé à une distance de la bride de tube (208) et supporté sur la bride (208) de manière à ce que, lors de l'insertion du corps (16) dans le tube (200), l'extrémité proximale du corps (16) vienne se mettre en contact avec le collier proximal supérieur (210), provoquant la flexion du collier proximal supérieur (210) qui quitte sa position de repos et à ce que, lorsque l'extrémité proximale (14) du corps (16) franchit le collier proximal supérieur (210), le collier proximal supérieur (210) soit suffisamment élastique pour retrouver sa position de repos. 35

2. Dispositif médical selon la revendication 1, dans lequel le collier proximal supérieur (210) est supporté sur le tube (200) par une pluralité de supports (212). 40
3. Dispositif médical selon les revendications 1 ou 2, dans lequel le collier proximal supérieur (210) a généralement la forme d'un polygone présentant un bord (214) qui s'incline vers l'intérieur le long d'une surface supérieure. 45
4. Dispositif médical selon la revendication 3, dans lequel le corps (16) est pourvu d'une bride (18) prévue autour de l'extrémité proximale (14) et le bord (214), incliné vers l'intérieur, de la surface interne du collier proximal supérieur (210) se termine au niveau d'un épaulement (218) pourvu d'un bord avant (218a) qui s'étend au-delà d'un point au niveau duquel une surface courbe de la bride (18) du corps se transforme en une surface plane, de manière à ce que le bord avant ne puisse pas se mettre facilement en contact avec la partie courbe de la bride de corps afin d'empêcher un retrait facile du tube. 50
5. Dispositif médical selon les revendications 1, 2, 3 ou 4, dans lequel le bord (134) du collier (132) possède une surface de butée plate (136a) et une surface inclinée (138) faisant face au côté distal, de manière à ce que, lorsque la protection (42) est poussée contre le tube (200), l'extrémité proximale (52) se déplace le long de la surface inclinée (138). 55

FIG-1



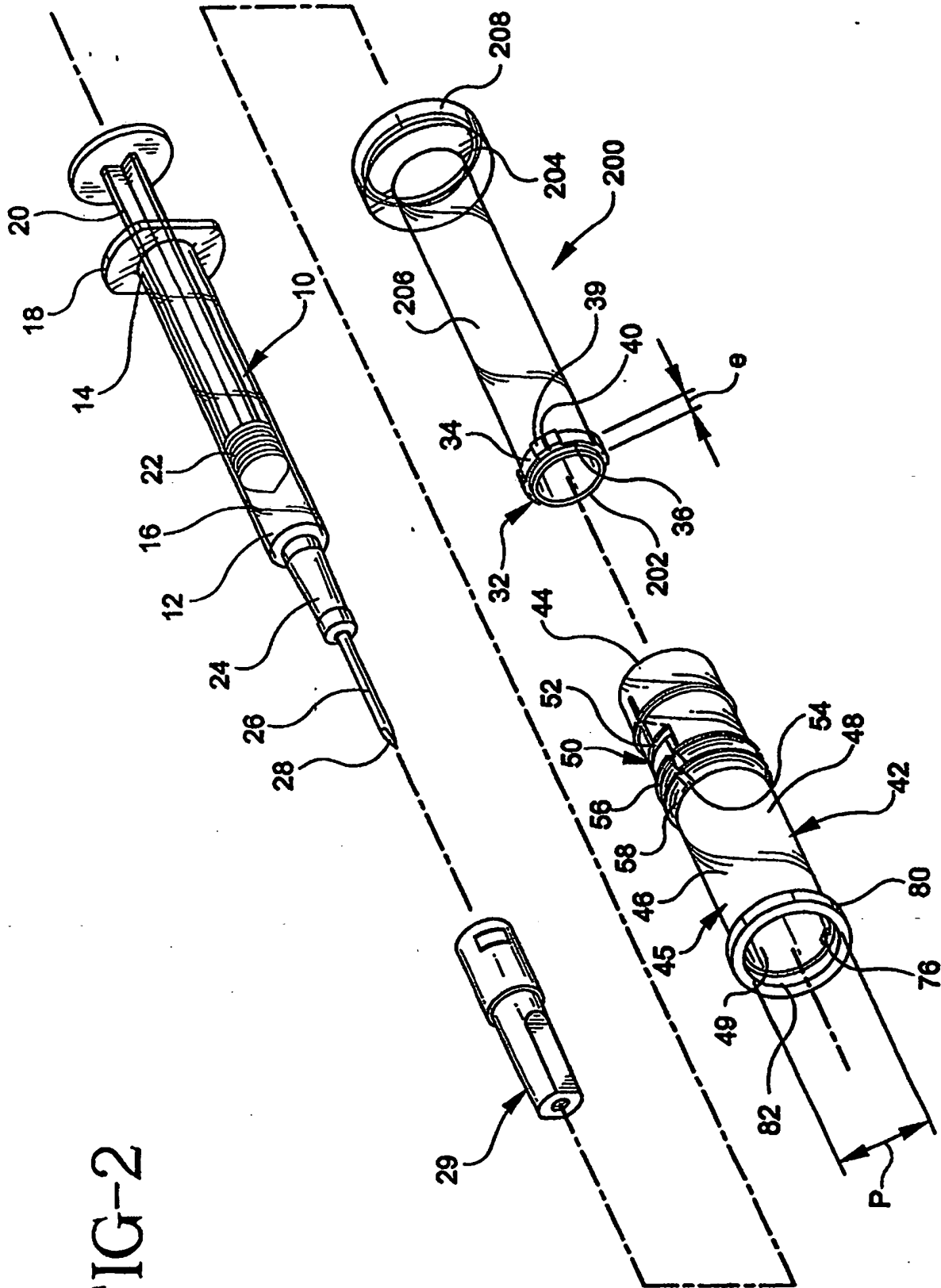


FIG-2

FIG-3

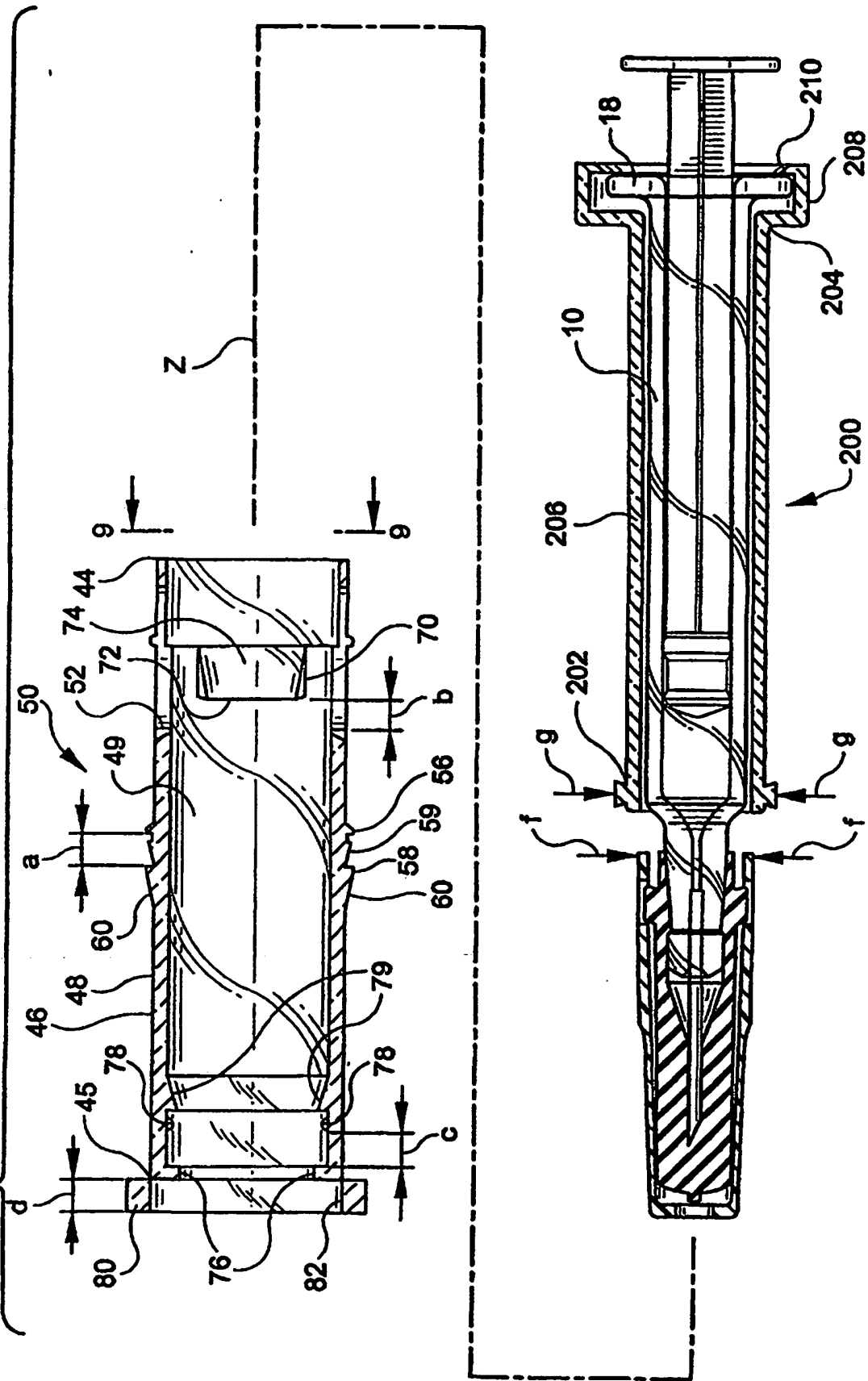


FIG-4

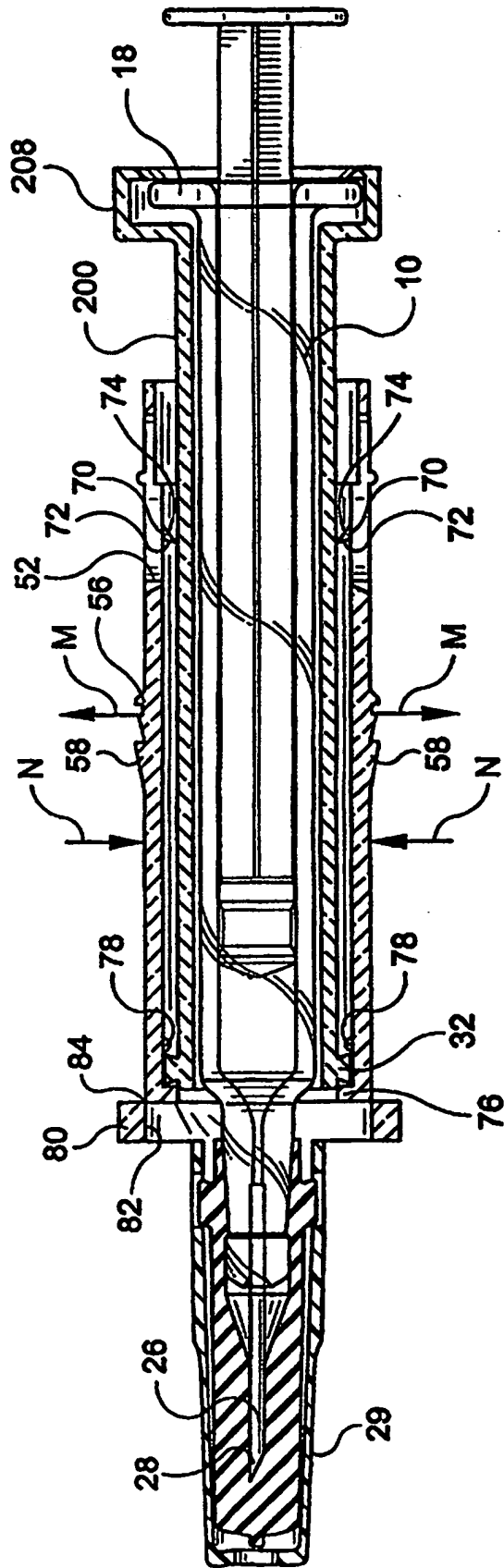


FIG-5

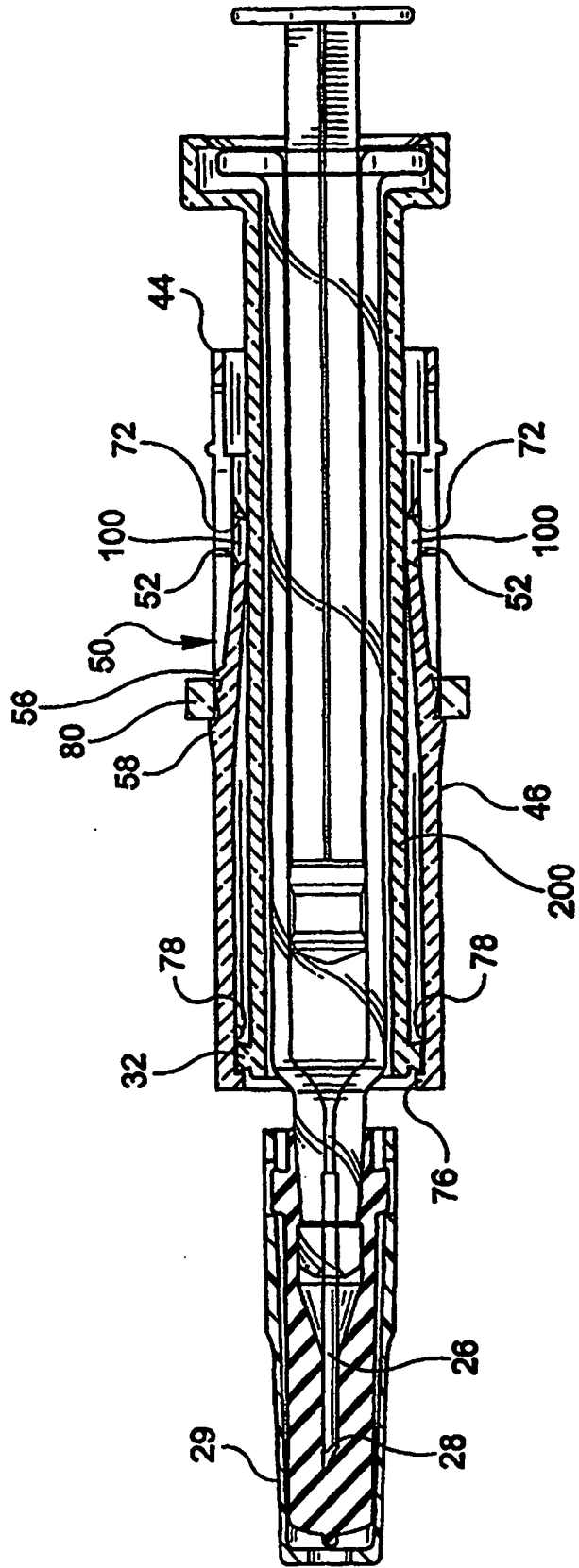


FIG-6

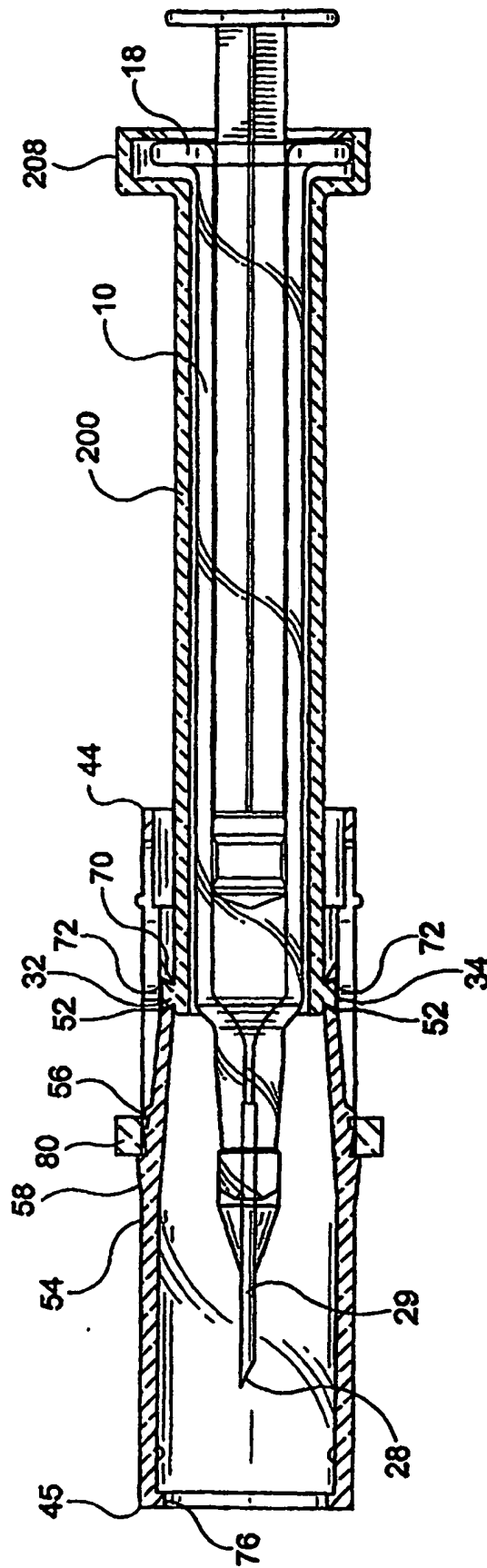


FIG-7

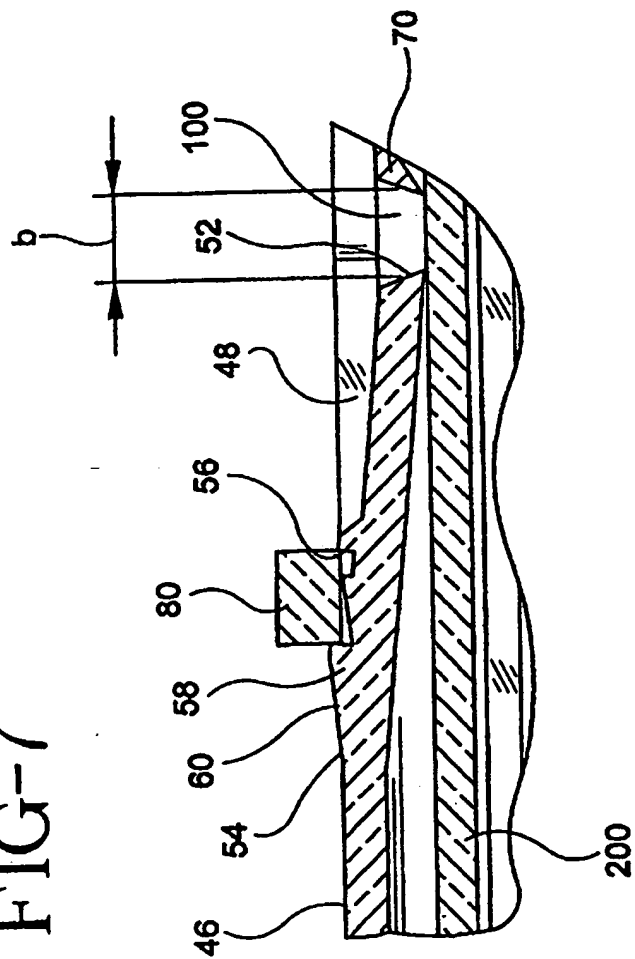
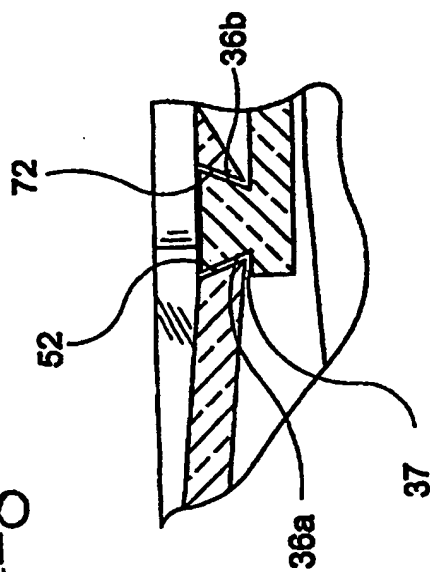


FIG-8



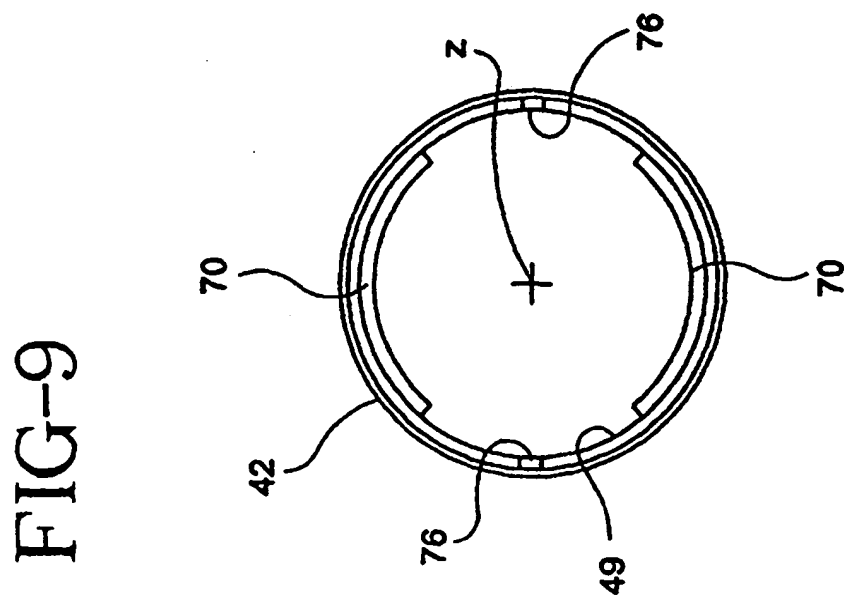
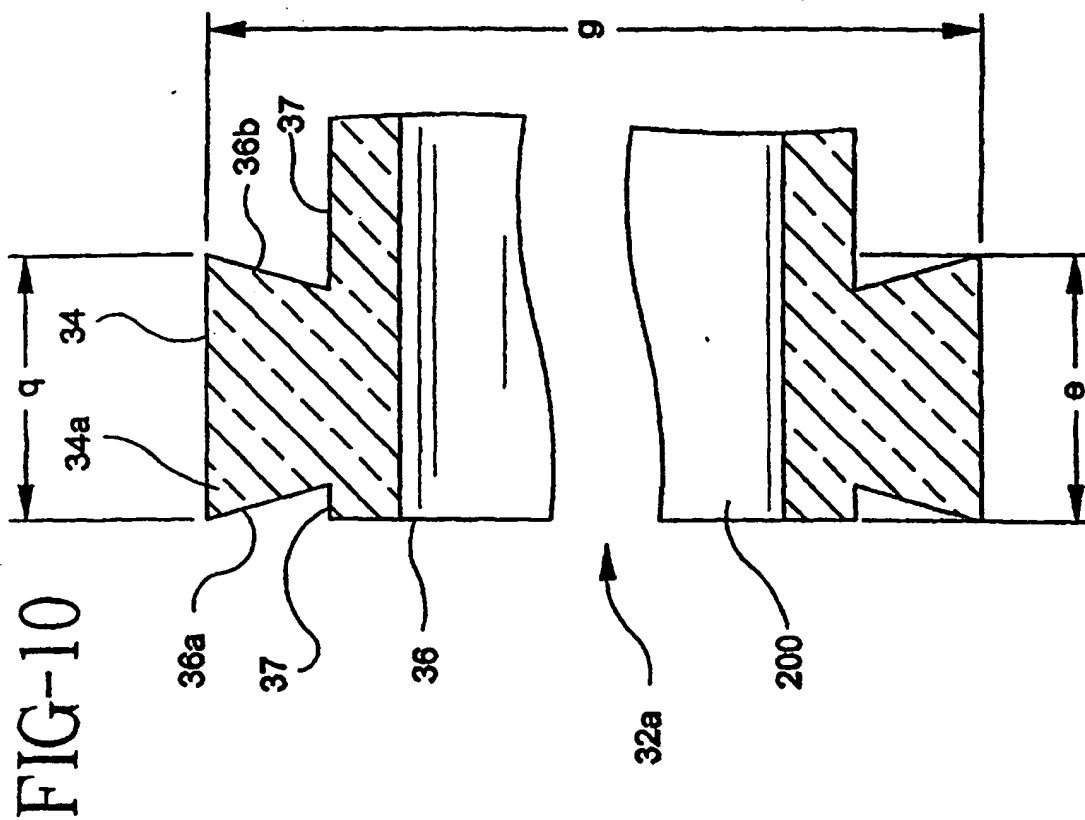


FIG-11

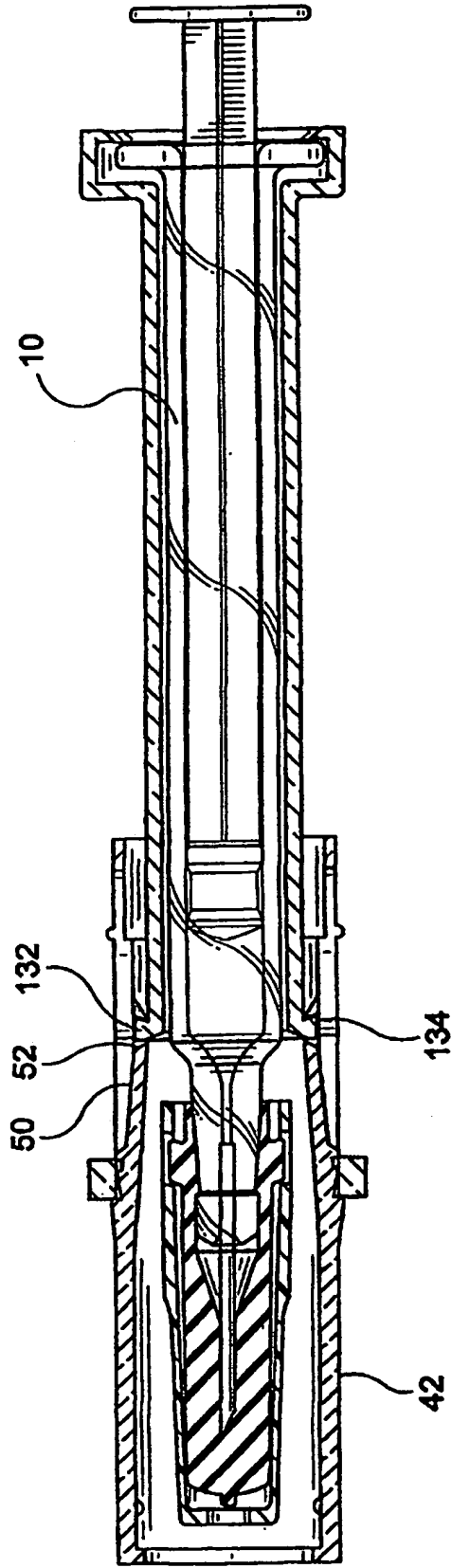


FIG-12

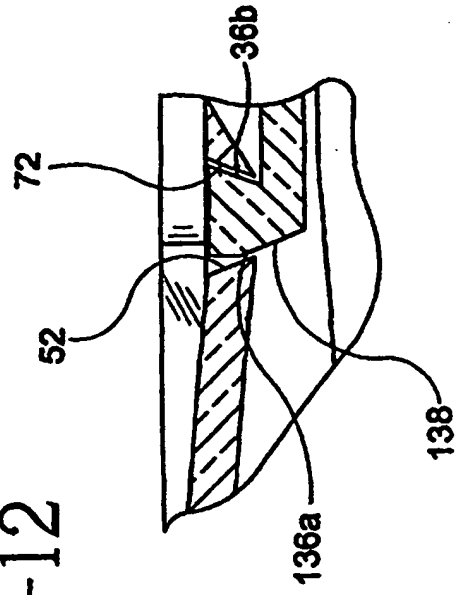


FIG-13

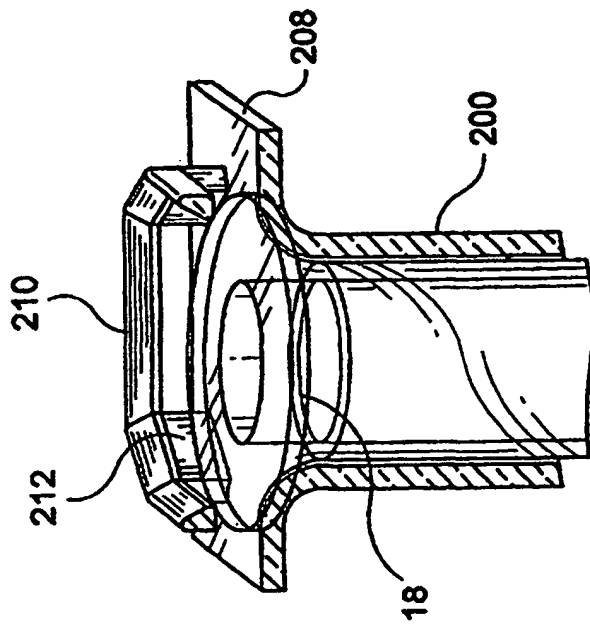


FIG-14

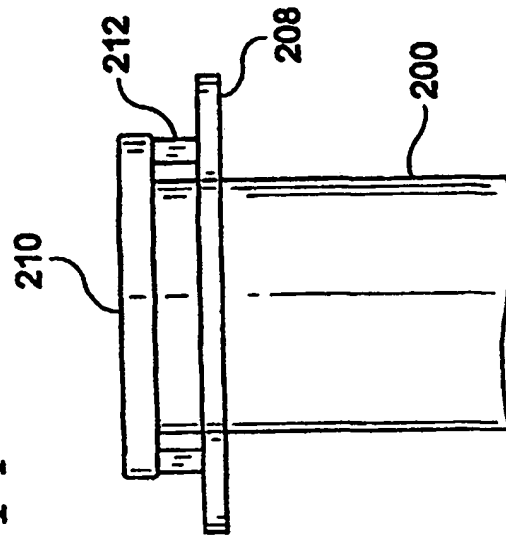


FIG-15

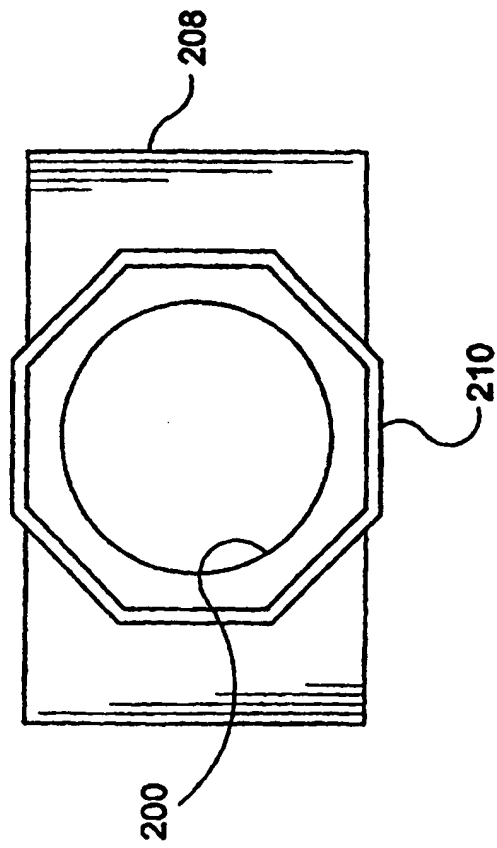


FIG-16

